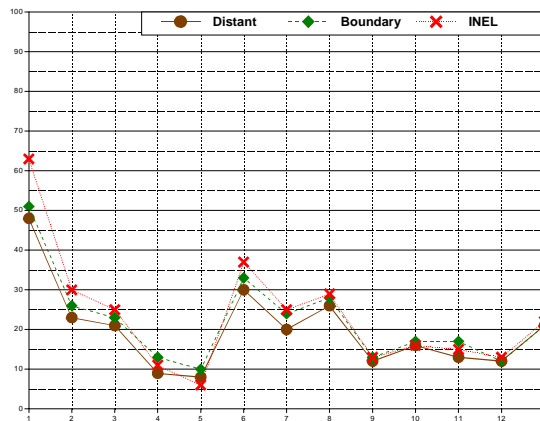


# INEL Offsite Environmental Surveillance Program Report: Second Quarter of 1996

Don Peterson  
Russell Mitchell  
Donny Roush



Environmental Science and Research Foundation  
February 1997



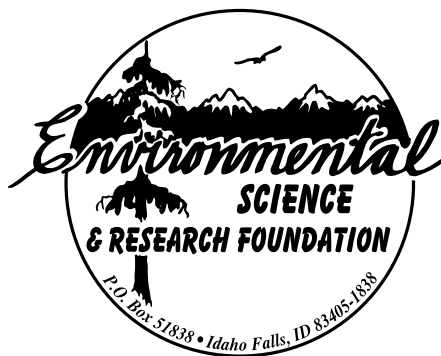
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# INEL Offsite Environmental Surveillance Program Report: Second Quarter of 1996

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**Environmental Science and Research Foundation**  
Doyle Markham, Executive Director

February 1997



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Environmental Science and Research Foundation  
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## Executive Summary

The Environmental Science and Research Foundation conducts the Idaho National Engineering Laboratory (INEL) Offsite Environmental Surveillance Program. The Foundation's environmental surveillance program is designed to monitor the effects, if any, of Department of Energy (DOE) activities on the offsite environment, to collect data to confirm compliance with applicable environmental laws and regulations, and to observe any trends in environmental levels of radioactivity. This report for the second quarter of 1996 includes the results of analyses conducted on samples of air, water, milk, game, sheep, and environmental radiation. A total of 526 samples were collected and analyzed. All concentrations of radioactivity found in these samples were consistent with historical levels. No evidence of radionuclides from the INEL was found in offsite samples. Concentrations of radionuclides found in all samples were below the guidelines set by both the Department of Energy and the Environmental Protection Agency (EPA) for radiation protection of the public.

## Program Description

The Foundation collected filters weekly from low-volume air samplers at 11 offsite locations. Four were at distant locations and seven at INEL boundary locations. An additional three samplers were operated on the INEL. Weekly measurements were made of gross alpha and gross beta concentrations in airborne particulates. Charcoal cartridges were screened weekly for the presence of Iodine-131. At the end of the quarter, weekly filters from each location were combined to form a composite sample for each location. These composites were then analyzed for gamma-emitting radionuclides. Selected composites were also submitted for Strontium-90 and transuranic analyses (Plutonium-238, Plutonium-239/240, and Americium-241).

Atmospheric moisture and precipitation samples were collected to monitor for tritium. Atmospheric moisture samples were collected for a period of approximately eight weeks. The Foundation collected two precipitation samples monthly, one onsite and one offsite, as well as a weekly onsite sample.

Drinking water samples were collected from 13 offsite locations and surface water samples were obtained from five sites. All water samples were analyzed for gross alpha, gross beta, and tritium concentrations.

The Foundation collected a weekly milk sample from a dairy in Idaho Falls and collected monthly milk samples from eight additional dairies around the INEL. All milk samples were analyzed for Iodine-131. Selected samples were also analyzed for either tritium or Strontium-90.

Annual sheep samples from flocks grazing on the INEL and in an area distant from the INEL were collected and analyzed for gamma-emitting radionuclides. Five road-killed game animals were also sampled and analyzed for gamma-emitting radionuclides.

The Foundation collected 13 offsite TLDs (environmental radiation dosimeters) that are analyzed semiannually to determine environmental radiation levels around the INEL.

## Summary of Second Quarter 1996 Results

During the second quarter of 1996, gross alpha and gross beta concentrations in low-volume air samples were within the expected range of values for background radioactivity. Mean concentrations of both gross alpha and gross beta were similar at onsite, distant, and boundary locations. Iodine-131 was not found in any air sample. No Cesium-137, Strontium-90, or transuranics (Americium-241, Plutonium-238, 239/240) were detected at any location.

Tritium was detected in one of two atmospheric moisture samples. The tritium, detected in a sample from a distant station, was attributed to historic, above-ground nuclear weapons tests and natural atmospheric processes.

Tritium was found in three of six precipitation samples. These samples were from both onsite and offsite (distant) locations. The tritium was attributed to natural atmospheric processes and historic nuclear weapons tests.

Tritium was detected in two of the 18 offsite water samples. Environmental tritium comes from natural processes and historic nuclear weapons tests. None of the samples contained detectable levels of gross alpha. Almost half of the samples contained concentrations of gross beta just above the minimum detectable concentration. Gross beta radioactivity in these water samples was attributed to naturally-occurring radionuclides in the earth's crust.

None of the milk samples collected during the second quarter contained detectable concentrations of Iodine-131. One of four samples had a detectable concentration of tritium; all five samples analyzed for Strontium-90 had detectable concentrations. The presence of these radionuclides was attributed to fallout from historic nuclear weapons tests.

Six sheep were analyzed for man-made gamma-emitting radionuclides; four of the six sheep grazed at onsite locations. Some of the onsite sheep had detectable concentrations of Cesium-137; one onsite sheep had a detectable concentration of Cobalt-60. None of the thyroids from the six sheep contained detectable concentrations of Iodine-131. Tissues from four of the five road-killed game animals contained detectable concentrations of Cesium-137. One of these four animals also had measurable concentrations of Cobalt-60. These radionuclides were attributed to either fallout from nuclear weapons testing or to ingestion of onsite soils contaminated with Cesium-137 from previous INEL operations.

The environmental radiation results were consistent with previously reported data and indicated no increase in detectable levels of environmental radiation due to INEL activities.

# Helpful Information for Readers

## Radionuclide Nomenclature

Radionuclides are sometimes expressed with the one- or two-letter chemical symbol for the element. A radionuclide is an unstable, or radioactive, form of an element. A given element may have many different radionuclides. Each is designated by a superscript number to the left of the chemical symbol. This number is the atomic weight of the radionuclide, equal to the number of protons and neutrons in its nucleus. Radionuclides which may be used in this report are shown in the following table:

<u>Symbol</u>	<u>Radionuclide</u>	<u>Symbol</u>	<u>Radionuclide</u>
<sup>3</sup> H	Tritium	<sup>131</sup> I	Iodine-131
<sup>7</sup> Be	Beryllium-7	<sup>134</sup> Cs	Cesium-134
<sup>51</sup> Cr	Chromium-51	<sup>137</sup> Cs	Cesium-137
<sup>54</sup> Mn	Manganese-54	<sup>144</sup> Ce	Cerium-144
<sup>58</sup> Co	Cobalt-58	<sup>181</sup> Hf	Hafnium-181
<sup>60</sup> Co	Cobalt-60	<sup>238</sup> Pu	Plutonium-238
<sup>65</sup> Zn	Zinc-65	<sup>239/240</sup> Pu	Plutonium-239/240
<sup>90</sup> Sr	Strontium-90	<sup>241</sup> Am	Americium-241
<sup>95</sup> Nb	Niobium-95		

## Scientific Notation

Scientific notation is used to express numbers which are very small and very large. A very small number will be expressed with a negative exponent, e.g.,  $1.3 \times 10^{-6}$ . To convert this number to the more commonly used form, the decimal point must be moved left by a number of places equal to the exponent (in this case, six). The number thus becomes 0.0000013.

For large numbers, those with a positive exponent, the decimal point is moved to the right by the number of places equal to the exponent. The number 1,000,000 (or one million) can be written as  $1.0 \times 10^6$ .

## Unit Prefixes

Units for very small and very large numbers are commonly expressed with a prefix. One example is the prefix *kilo*, abbreviated k, which means 1,000 of a given unit. A kilometer is therefore equal to 1,000 meters. Prefixes that may be used in this report are:

<u>Prefix</u>	<u>Abbreviation</u>	<u>Meaning</u>
milli	m	1/1,000 (= $1 \times 10^{-3}$ )
micro	$\mu$	1/1,000,000 (= $1 \times 10^{-6}$ )
pico	p	1/1,000,000,000,000 (= $1 \times 10^{-12}$ )

## Units of Radioactivity and Radiation Exposure and Dose

The basic unit of radioactivity used in this report is the curie, abbreviated Ci. The curie is defined as the amount of radioactivity equivalent to 37 billion nuclear transformations per second. Historically, this was based upon the radioactivity from one gram of the radionuclide Radium-226. For any other radionuclide, one curie is the amount of that radionuclide that decays at this same rate.

Radiation exposure is expressed in terms of the Roentgen (R), the amount of ionization produced by gamma radiation in air. Dose is given in units of "Roentgen equivalent man," or "rem," which takes into account the effect of radiation on tissues. For the types of environmental radiation generally encountered, the unit of Roentgen is approximately numerically equal to the unit of rem.

## **Units of Environmental Concentrations**

Concentration of radioactivity in air and milk samples is expressed in units of microcuries per milliliter ( $\mu\text{Ci/mL}$ ). Concentrations in water samples are expressed as picocuries per liter ( $\text{pCi/l}$ ); federal standards are expressed in these units. Radioactivity in foodstuffs are given in microcuries per gram ( $\mu\text{Ci/g}$ ), dry weight. Radioactivity in soil samples is expressed as picocuries per gram ( $\text{pCi/g}$ ), dry weight. Annual human radiation exposure, measured by environmental dosimeters, is expressed in units of milliRoentgens (mR). This is sometimes expressed in terms of dose as millirem (mrem). Not all of the above sample types may appear in this particular report.

## **Uncertainty of Measurements**

Due to many variables, there is always an uncertainty associated with the measurement of environmental contaminants. For radioactivity, the predominant source of uncertainty is due to the inherent statistical nature of radioactive decay events, particularly at the low activity levels encountered in environmental samples. The uncertainty of a measurement is denoted by following the result with a " $\pm$ " (uncertainty) term. This report follows convention in reporting the uncertainty as a 95% confidence limit (or interval), designated in the tables as " $\pm 2s$ ." That means there is approximately a 95% level of confidence that the real concentration in the sample lies somewhere between the measured (reported) concentration minus the uncertainty term and the measured (reported) concentration plus the uncertainty term.

## **Negative Numbers as Results**

Environmental measurements are frequently conducted at levels where the contaminant, such as radioactivity, cannot be distinguished from natural background levels. In this case, the result will still be reported by the analytical laboratory, even though it is below the measurement system's approximate minimum detectable concentration, or is less than zero. Negative values occur when the measured result is less than a pre-established average background level for the particular system and procedure used. These values, rather than "not detectable" or "zero," are reported to better enable statistical analyses and to observe trends in the data.

## **Gross versus Specific Analyses**

Many of the radiological analyses of environmental samples yield information only about the overall, or gross, amount of a particular type of radiation (e.g., gross beta), rather than identifying and quantifying specific radionuclides. For example, rather than performing an analysis for particular gamma-emitting radionuclides, called gamma spectroscopy, one can do a gross gamma or, more commonly, a gross beta analysis, since gamma-emitting radionuclides also emit beta particles. This type of analysis is an effective screening tool and is much quicker and less costly than specific radionuclide analyses.



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### 1. Introduction

Consistent with requirements of applicable Department of Energy (DOE) Orders, the Foundation's environmental surveillance program is designed to monitor the effects, if any, of DOE activities on the offsite environment, to collect data which verifies compliance with applicable environmental laws and regulations, and to observe trends in environmental levels of radioactivity. This work is performed under DOE Contract DE-AC07-ID13268.

This quarterly report summarizes the data collected by the Foundation's INEL Offsite Environmental Surveillance Program during the period April 1 through June 30, 1996. The scope of the Foundation's sampling program is outlined in Table 1. Most analyses for the surveillance program were performed by Idaho State University's Environmental Monitoring Laboratory. Other analyses were performed by Quanterra Laboratory, a commercial laboratory located in Richland, Washington.

A large portion of environmental results are near background levels of radioactivity; many results are near the detection limits of the laboratory procedures. Table 2 summarizes the approximate minimum detectable concentrations (MDC) of radioactivity that the laboratories can detect and quantify for a given sample type and analysis. All results are reported with an associated 2s ("two sigma") uncertainty term. The Foundation has adopted the following method for interpreting analytical results near the minimum detectable concentration. Results less than or equal to the 2s uncertainty term, which includes some results that are negative, are considered as "not detected." For results greater than 2s (the 95% confidence level), but not exceeding 3s (the 99% confidence interval), detection of the radioactivity is questionable. These results may exceed the 2s level simply due to random statistical fluctuations. This is expected to occur approximately 2.5% of the time. Results exceeding 3s are interpreted as indicating that radioactivity was detected.

Where appropriate, the results in this report are compared to the following:

- ▶ For air, concentrations are compared to the DOE Derived Concentration Guides. This is the concentration of a radionuclide that, under conditions of continuous exposure, would result in an effective dose equivalent of 100 mrem (the DOE standard for members of the public);
- ▶ For drinking water, concentrations are compared to the Environmental Protection Agency's Maximum Contaminant Level. This is the maximum permissible level of a contaminant in water that is delivered to any user of a community water system.

# 1. Introduction

Table 1 Summary of the Foundation's Environmental Surveillance Program				
Sample Type Analysis	Collection Frequency	Locations		
		Distant	Boundary	INEL
<b>Air</b>				
Gross Alpha	weekly	Blackfoot, Craters of the Moon, Idaho Falls, Rexburg	Arco, Atomic City, FAA Tower, Howe, Monteview, Mud Lake, Reno Ranch	Main Gate, EFS, Van Buren
Gross Beta <sup>131</sup> I	weekly	Blackfoot, Craters of the Moon, Idaho Falls, Rexburg	Arco, Atomic City, FAA Tower, Howe, Monteview, Mud Lake, Reno Ranch	Main Gate, EFS, Van Buren
Gamma Spec Particulate Mass	quarterly	Blackfoot, Craters of the Moon, Idaho Falls, Rexburg	Arco, Atomic City, FAA Tower, Howe, Monteview, Mud Lake, Reno Ranch	Main Gate, EFS, Van Buren
<sup>90</sup> Sr Transuranics	quarterly	Rotating schedule	Rotating schedule	Rotating schedule
<b>Air Moisture</b>				
Tritium	4 to 13 weeks	Idaho Falls	Atomic City	None
<b>Precipitation</b>				
Tritium	monthly	Idaho Falls	None	CFA
Tritium	weekly	None	None	EFS
<b>Surface H<sub>2</sub>O</b>				
Gross Alpha, Gross Beta, <sup>3</sup> H	quarterly → semiannually →	Twin Falls, Buhl, Hagerman Idaho Falls, Bliss	None	None
<b>Drinking H<sub>2</sub>O</b>				
Gross Alpha Gross Beta, <sup>3</sup> H	semiannually	Aberdeen, Blackfoot, Carey, Idaho Falls, Fort Hall, Minidoka, Roberts, Shoshone	Arco, Atomic City, Howe, Monteview, Mud Lake, Reno Ranch	None
<b>Milk</b>				
<sup>131</sup> I	weekly	Idaho Falls	None	None
<sup>131</sup> I	monthly	Blackfoot, Carey, Dietrich, Minidoka, Roberts	Howe, Terreton, Arco	None
Tritium <sup>90</sup> Sr	annually	Blackfoot, Carey, Dietrich, Idaho Falls, Minidoka, Roberts	Howe, Terreton, Arco	None
<b>Potatoes</b>				
Gamma Spec <sup>90</sup> Sr	annually	Blackfoot, Idaho Falls, Rupert	Arco, Mud Lake	None
<b>Wheat</b>				
Gamma Spec <sup>90</sup> Sr	annually	American Falls, Blackfoot, Dietrich, Idaho Falls, Minidoka, Carey	Arco, Monteview, Mud Lake, Tabor, Terreton	None
<b>Lettuce</b>				
Gamma Spec <sup>90</sup> Sr	annually	Blackfoot, Carey, Idaho Falls, Pocatello	Arco, Atomic City, Howe, Mud Lake	None
<b>Fish</b>				
Gamma Spec	annually	None	None	Big Lost River
<b>Sheep</b>				
Gamma Spec	annually	Blackfoot	None	INEL grazing areas
<b>Waterfowl</b>				
Gamma Spec <sup>90</sup> Sr Transuranics	annually	Fort Hall	None	Waste disposal ponds
<b>Game</b>				
Gamma Spec	varies	None	None	INEL roads
<b>Soil</b>				
Gamma Spec <sup>90</sup> Sr Transuranics	biennially	Carey, Crystal Ice Caves, Blackfoot, St. Anthony	Butte City, Monteview, Atomic City, FAA Tower, Howe, Mud Lake (2), Reno Ranch	None
<b>TLDs</b>				
Gamma Radiation	semiannual	Aberdeen, Blackfoot, Craters of the Moon, Idaho Falls, Minidoka, Rexburg, Roberts	Arco, Atomic City, Howe, Monteview, Mud Lake, Reno Ranch	None

## 1. Introduction

**Table 2**  
**Summary of Approximate Minimum Detectable Concentrations for Radiological Analyses**  
**(Second Quarter 1996)**

<u>Sample Type</u>	<u>Analysis</u>	<u>Approximate Minimum Detectable Concentration<sup>a</sup> (MDC)</u>	<u>Derived Concentration Guide<sup>b</sup> (DCG)</u>	<u>Drinking Water Detection Limits<sup>c</sup></u>
Air (particulate filter) <sup>d</sup>	Gross alpha	1 x 10 <sup>-15</sup> μCi/ml	2 x 10 <sup>-14</sup> μCi/ml	--
	Gross beta	4 x 10 <sup>-15</sup> μCi/ml	3 x 10 <sup>-12</sup> μCi/ml	--
	Specific gamma ( <sup>137</sup> Cs)	2 x 10 <sup>-15</sup> μCi/ml	4 x 10 <sup>-10</sup> μCi/ml	--
	<sup>238</sup> Pu	2 x 10 <sup>-18</sup> μCi/ml	3 x 10 <sup>-14</sup> μCi/ml	--
	<sup>239/240</sup> Pu	3 x 10 <sup>-18</sup> μCi/ml	2 x 10 <sup>-14</sup> μCi/ml	--
	<sup>241</sup> Am	2 x 10 <sup>-18</sup> μCi/ml	2 x 10 <sup>-14</sup> μCi/ml	--
	<sup>90</sup> Sr	3 x 10 <sup>-17</sup> μCi/ml	9 x 10 <sup>-12</sup> μCi/ml	--
Air (charcoal cartridge) <sup>d</sup>	<sup>131</sup> I	4 x 10 <sup>-15</sup> μCi/ml	4 x 10 <sup>-10</sup> μCi/ml	--
Air (atmospheric moisture) <sup>e</sup>	<sup>3</sup> H	4 x 10 <sup>-12</sup> μCi/ml	1 x 10 <sup>-7</sup> μCi/ml	--
Air (precipitation)	<sup>3</sup> H	1 x 10 <sup>-7</sup> μCi/ml	2 x 10 <sup>-3</sup> μCi/ml	--
Water (drinking & surface)	Gross alpha	4 pCi/l	30 pCi/l	3 pCi/l
	Gross beta	2 pCi/l	100 pCi/l	4 pCi/l
	<sup>3</sup> H	100 pCi/l	2 x 10 <sup>6</sup> pCi/l	1000 pCi/l
Milk	<sup>131</sup> I	2 x 10 <sup>-9</sup> μCi/ml	--	--
	<sup>3</sup> H	1 x 10 <sup>-7</sup> μCi/ml	--	--
	<sup>90</sup> Sr	5 x 10 <sup>-10</sup> μCi/ml	--	--
Thyroid tissue	<sup>131</sup> I	3 x 10 <sup>-7</sup> μCi/g	--	--
Liver tissue	<sup>137</sup> Cs	5 x 10 <sup>-9</sup> μCi/g	--	--
Muscle tissue	<sup>137</sup> Cs	4 x 10 <sup>-9</sup> μCi/g	--	--
<p>a. The MDC is an estimate of the concentration of radioactivity in a given sample type that can be identified with a 95% level of confidence and a precision of plus or minus 100% under a specified set of typical laboratory measurement conditions.</p> <p>b. DCGs, set by the DOE, represent reference values for radiation exposure. They are based on a radiation dose of 100 mrem/yr for exposure through a particular exposure mode such as direct exposure, inhalation, or ingestion of water.</p> <p>c. These limits are required by the National Primary Drinking Water Regulations (40 CFR 141). The "detection limit" is the terminology used by the EPA and means the same as the MDC defined above.</p> <p>d. The approximate MDC is based on an average filtered air volume (pressure corrected) of 570 m<sup>3</sup>/week.</p> <p>e. The approximate MDC is expressed for tritium (as tritiated water) in air, and is based on an average filtered air volume of 25 m<sup>3</sup>, assuming an average sampling period of eight weeks.</p>				

## 2. Air Sampling

### 2.1 Sampling Methods

#### 2.1.1 Low-Volume Air Samplers

Airborne particulate radioactivity was continuously monitored by 14 air samplers (Figure 1), designed to provide an effective network to detect INEL releases of radioactivity. Four offsite air samplers were designated as distant, or background, stations and seven are designated as boundary stations. Three air samplers are situated on the INEL. Distant locations were used to make comparisons of airborne concentrations of radioactivity with boundary and onsite locations. As part of the quality assurance program, two replicate samplers, located in Mud Lake and near the INEL Main Gate, were operated adjacent to regular air samplers to provide a means of comparing data.

Each air sampler averaged a flow of approximately 50 l/min (2 ft<sup>3</sup>/min) through a filter head consisting of two types of filters—a 1.2-micrometer pore size particulate filter and a charcoal cartridge for the monitoring of radioactive iodine. Filters on each sampler were changed weekly. In order to be considered a valid sample, each filter must sample a pressure-corrected air volume of at least 200 m<sup>3</sup>, or about 7000 ft<sup>3</sup>. Filters sample an average air volume of about 570 m<sup>3</sup> (20,000 ft<sup>3</sup>).

Charcoal cartridges were screened in batches weekly for <sup>131</sup>I activity. If activity was detected in any batch that was greater than a preset action level, cartridges were then analyzed individually. Particulate filters were counted each week for gross (nonspecific) beta activity in a low-background beta counter after waiting a minimum of four days for the naturally occurring decay products of radon and thoron to decay. The particulate filters were also counted for gross alpha activity.

At the end of the quarter, weekly filters from each location were combined to form a composite. All composites were then analyzed by gamma spectrometry for specific radionuclides. Selected composites were also analyzed for <sup>90</sup>Sr or transuranic radionuclides (<sup>238</sup>Pu, <sup>239/240</sup>Pu, and <sup>241</sup>Am).

## 2. Air Sampling

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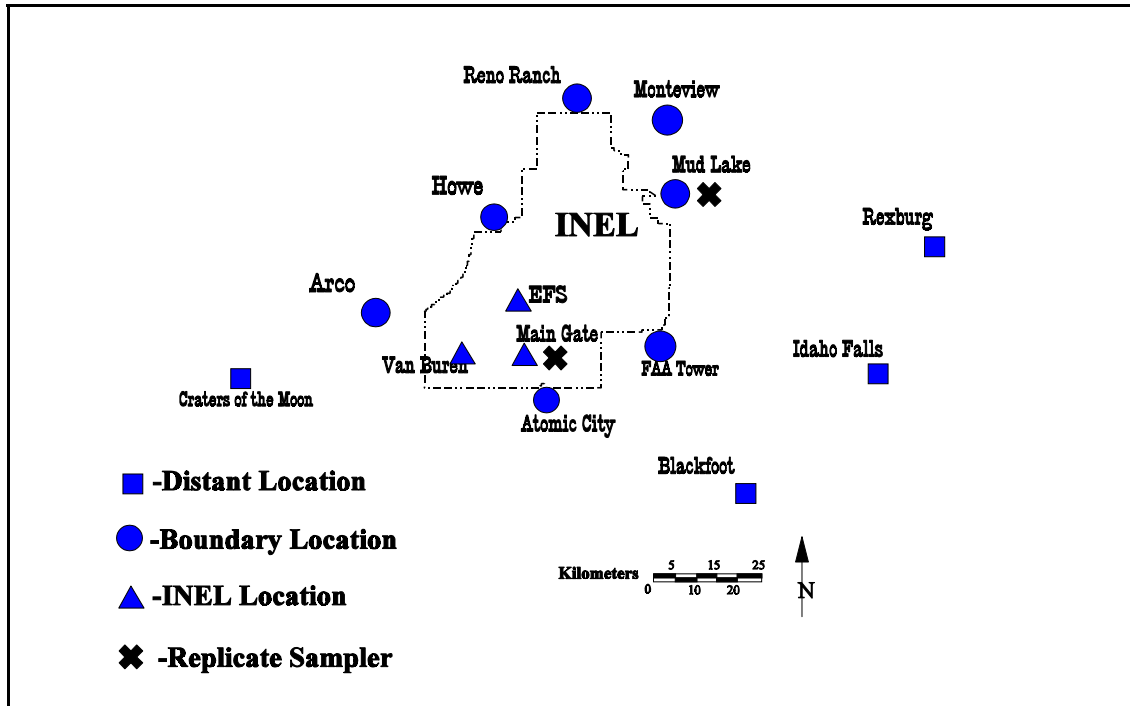


Figure 1 Air Sampling Location Map

### 2.1.2 Atmospheric Moisture Samplers

Two air samplers, located in Atomic City and Idaho Falls, collected atmospheric moisture for tritium analysis. Air was passed through a column of silica gel that absorbs water vapor in the air. Tritium concentrations were determined by extracting water from the silica gel and counting the water sample by liquid scintillation.

### 2.1.3 Precipitation Samplers

When available, weekly precipitation samples were collected at the Experimental Field Station (EFS) on the INEL. In addition, two samples were collected monthly: one at the Central Facilities Area on the INEL and one in Idaho Falls. All precipitation samples were analyzed for tritium by liquid scintillation.

## 2. Air Sampling

### 2.2 Results

#### 2.2.1 Low-Volume Air Samplers

No  $^{131}\text{I}$  was detected in any of the weekly charcoal cartridge batches analyzed during the second quarter, thus no analyses of individual cartridges were required. The minimum detectable concentration was approximately  $4 \times 10^{-15} \mu\text{Ci/ml}$ .

All gross alpha concentrations were within the expected range of background levels (data summaries are presented in Figure 2, Table 3, and Table A-1). Figure 2 indicates that the levels of airborne radioactivity for the three groups of stations (distant, boundary, and INEL) are similar, within uncertainty limits, and track each other closely over the 13 weeks. This is an indication that the fluctuations occurred over the entire sampling network and are therefore not likely caused by a localized source such as a facility at the INEL. Table 3 indicates the mean gross alpha concentration for Mud Lake and the replicate are slightly higher than for the other boundary stations. However, comparison with the distant stations indicates background levels of gross alpha were not exceeded at Mud Lake. The quarterly mean gross alpha concentrations for the onsite and boundary locations were not statistically higher than the mean for the distant locations (see Table 3).

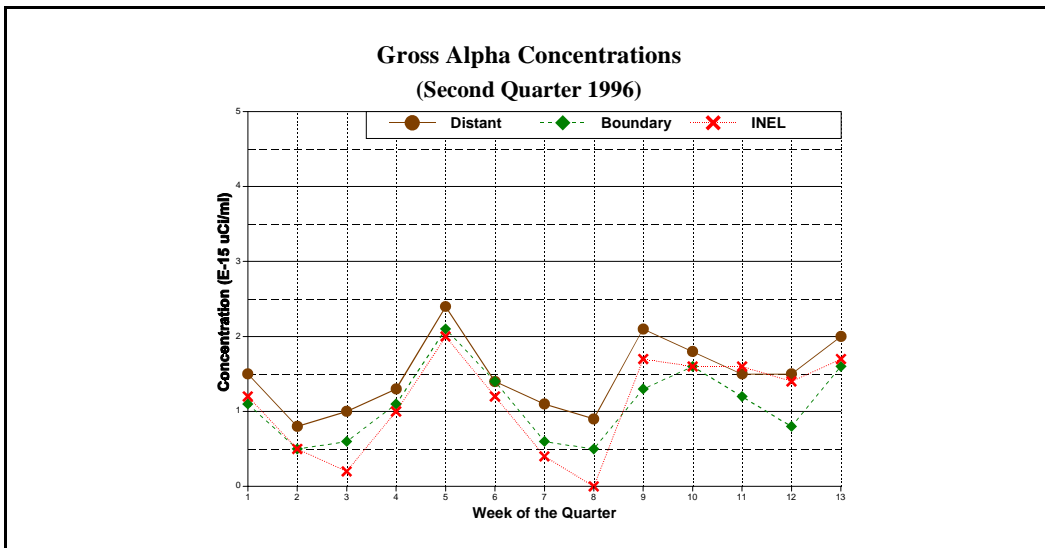


Figure 2 Weekly Gross Alpha Concentrations



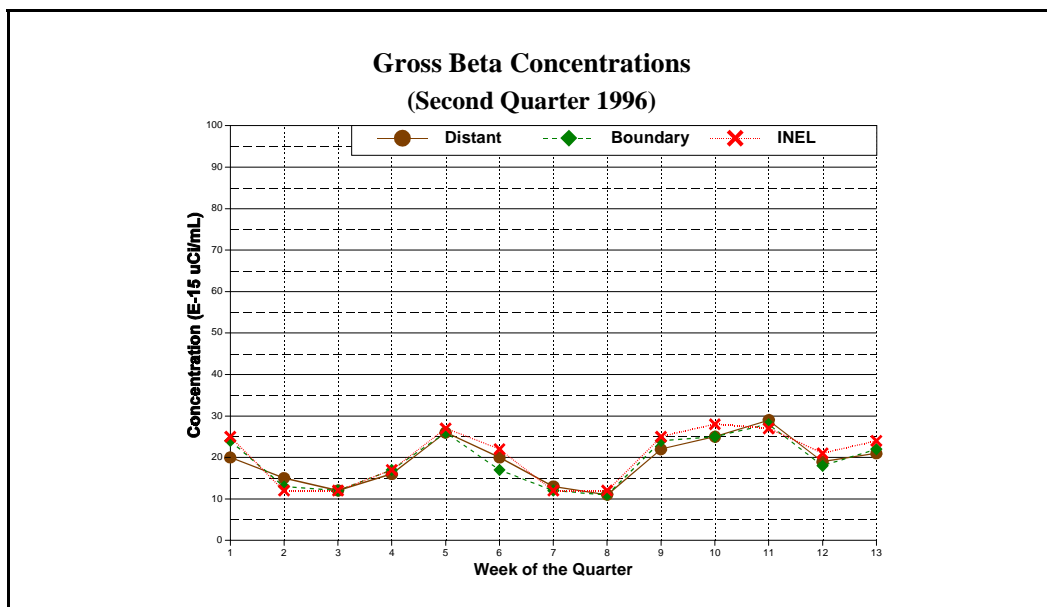
## 2. Air Sampling

<b>Table 3</b>				
<b>Gross Alpha Concentrations in Air</b>				
<b>(Second Quarter 1996)</b>				
<b>Group</b>	<b>Location</b>	<b>Number of Samples</b>	<b>Gross Alpha Concentration</b>	
			<b>Range of Samples</b>	<b>Mean with 95% Confidence Interval</b>
Distant	Blackfoot	13	0.7 - 3.1	1.8 ± 0.4
	Craters of the Moon	13	- 0.2 - 2.5	1.2 ± 0.4
	Idaho Falls	13	0.5 - 2.2	1.4 ± 0.3
	Rexburg	13	0.7 - 2.4	1.5 ± 0.4
			<b>Group Mean</b>	<b>1.5 ± 0.2</b>
Boundary	Arco	13	0.2 - 2.2	1.0 ± 0.3
	Atomic City	13	0.5 - 1.8	1.0 ± 0.2
	FAA Tower	13	0.2 - 2.0	1.2 ± 0.3
	Howe	13	-0.5 - 2.6	1.0 ± 0.5
	Montevieu	13	0.1 - 2.8	1.2 ± 0.5
	Mud Lake (Replicate)	13 (13)	0.4 - 2.4 (0.2 - 3.4)	1.5 ± 0.4 (1.5 ± 0.5)
	Reno Ranch	13	0.2 - 1.7	0.9 ± 0.3
			<b>Group Mean</b>	<b>1.1 ± 0.1</b>
INEL	EFS	13	0.1 - 1.8	1.1 ± 0.3
	Main Gate (Replicate)	13 (13)	-0.6 - 2.7 (0.1 - 2.0)	1.2 ± 0.6 (0.9 ± 0.3)
	Van Buren	13	0.2 - 1.9	1.1 ± 0.3
			<b>Group Mean</b>	<b>1.1 ± 0.2</b>
DOE Derived Concentration Guide				20

All gross beta concentrations were also within the expected range of background levels (data summaries are presented in Figure 3, Table 4, and Table A-2). Figure 3 indicates that the levels of airborne radioactivity for the three groups of stations (distant, boundary, and INEL) are similar, within uncertainty limits, and closely track each other over the 13 weeks. This is an indication that the fluctuations occurred over the entire sampling network and are therefore not likely caused by a localized source such as a facility at the INEL. Quarterly means of gross beta concentrations for the onsite and boundary locations were not statistically higher than the mean for the distant locations (see Table 4).

The gross alpha and gross beta data for the Mud Lake and Main Gate quality assurance replicates assisted in data validation. The Mud Lake and Main Gate mean values were not statistically different from their respective replicate mean values (see Tables 3 and 4), as determined by two-tailed Student's t-tests. Linear correlation tests indicated the Mud Lake and Main Gate weekly gross beta results

## 2. Air Sampling



**Figure 3 Weekly Gross Beta Concentrations**

correlated well at the 95% confidence level with their respective replicate results<sup>1</sup>. Correlation tests were performed only for those gross alpha data greater than their respective  $2s$  uncertainty terms. Correlation was poor, particularly for the Mud Lake station<sup>1</sup>. This is attributed to the detected levels of gross alpha being close to the minimum detectable concentration (see Table 2).

Quarterly composite samples from all sampling stations were analyzed for gamma-emitting radionuclides. Beryllium-7, a naturally-occurring gamma-emitting radionuclide produced by cosmic rays in the atmosphere, was detected in all of the composites. Cesium-137, sometimes detected on composite samples and whose presence is generally attributed to fallout from nuclear weapons, was not detected in any of the samples.

Several composite samples were also selected for analysis of  $^{90}\text{Sr}$  and transuranics of interest ( $^{241}\text{Am}$ ,  $^{238}\text{Pu}$ , and  $^{239/240}\text{Pu}$ ). No detectable concentrations of  $^{90}\text{Sr}$  or transuranics were detected at any of the locations.

<sup>1</sup> For Mud Lake,  $r_{\alpha} = 0.16$  ( $n = 9$ ),  $r_{\beta} = 0.89$  ( $n = 13$ ); for the Main Gate,  $r_{\alpha} = 0.45$  ( $n = 6$ ) and  $r_{\beta} = 0.90$  ( $n = 13$ ).

## 2. Air Sampling

<b>Table 4</b>				
<b>Gross Beta Concentrations in Air</b>				
<b>(Second Quarter 1996)</b>				
<b>Group</b>	<b>Location</b>	<b>Number of Samples</b>	<b>Gross Beta Concentration</b>	
			<b>Range of Samples</b>	<b>Mean with 95% Confidence Interval</b>
Distant	Blackfoot	13	8 - 27	18 ± 4
	Craters of the Moon	13	11 - 38	23 ± 4
	Idaho Falls	13	10 - 22	16 ± 2
	Rexburg	13	9 - 32	20 ± 4
	<b>Group Mean</b>			<b>19 ± 2</b>
Boundary	Arco	13	10 - 26	18 ± 3
	Atomic City	13	6 - 30	20 ± 4
	FAA Tower	13	8 - 30	18 ± 4
	Howe	13	11 - 27	20 ± 3
	Monteviu	13	9 - 26	18 ± 4
	Mud Lake (Replicate)	13 (13)	10 - 32 (12 - 35)	21 ± 4 (23 ± 5)
	Reno Ranch	13	9 - 27	19 ± 4
<b>Group Mean</b>			<b>19 ± 1</b>	
INEL	EFS	13	11 - 31	21 ± 4
	Main Gate (Replicate)	13 (13)	11 - 33 (12 - 28)	23 ± 5 (19 ± 3)
	Van Buren	13	7 - 28	18 ± 4
<b>Group Mean</b>			<b>20 ± 2</b>	
DOE Derived Concentration Guide				3000

### 2.2.2 Atmospheric Moisture Samplers

Atmospheric moisture samples were obtained from Idaho Falls and Atomic City in April 1996, representing moisture collected between February and April. The sample from Idaho Falls contained detectable tritium at  $(8.1 \pm 6.4) \times 10^{-14}$   $\mu\text{Ci/ml}$ . As of this writing, data from other samples taken during this report period were not available. Atmospheric tritium detected at stations distant from the INEL, such as Idaho Falls, is likely due to both natural sources (cosmic ray interactions in the atmosphere) and man-made sources (historic nuclear weapons testing).

## 2. Air Sampling

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### 2.2.3 Precipitation Samplers

Six precipitation samples were collected in the second quarter and analyzed for tritium. Tritium was detected in three samples, one from each of the sampling locations: Idaho Falls offsite; CFA (Central Facilities Area) and EFS (Experimental Field Station) onsite. The highest concentration was at the Idaho Falls location:  $(3.6 \pm 1.0) \times 10^{-7}$   $\mu\text{Ci/ml}$ , collected in April. For CFA the concentration was  $(1.3 \pm 1.0) \times 10^{-7}$   $\mu\text{Ci/ml}$ , collected in April. For EFS the concentration was  $(1.8 \pm 1.0) \times 10^{-7}$   $\mu\text{Ci/ml}$ , collected in mid-May. A reanalysis was performed for the Idaho Falls sample. The result was still slightly positive, but closer to levels detected in the other two samples:  $(1.1 \pm 1.0) \times 10^{-7}$   $\mu\text{Ci/ml}$ . Although tritium attributable to airborne releases from ICPP operations at the INEL was found in several onsite precipitation samples a few years ago, the similar concentrations found at distant and INEL locations indicate this quarter's results are more likely due to environmental tritium from natural atmospheric processes and historic nuclear weapons testing.

**Russ Mitchell, environmental scientist, prepares an instrument at EFS (Experimental Field Station) to record and measure precipitation events.**

### 3. Water Sampling

#### 3.1 Methods

Water samples were collected in early May from 13 drinking water locations and five surface water locations, four in the Magic Valley area and one from Idaho Falls (see Figure 4). Drinking water sampling locations were local businesses. Surface water locations included three springs in the Thousand Springs area (Figure 4). These springs are some of the outlets for the Snake River Plain Aquifer, which flows beneath the INEL. Each water sample was analyzed for gross (nonspecific) alpha and gross beta activity by evaporating a portion of the sample on a stainless steel plate and counting the residue. Tritium concentrations were determined by analyzing samples using liquid scintillation.

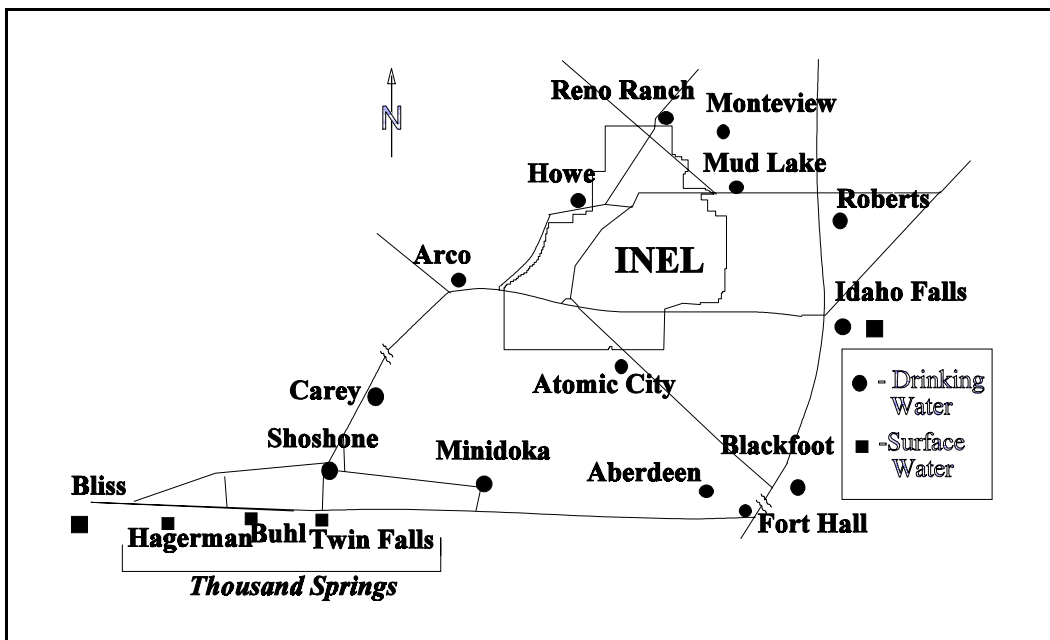


Figure 4 Water Sampling Locations

### 3. Water Sampling

#### 3.2 Results

Two of the drinking water samples (Blackfoot and Fort Hall) showed measurable concentrations of  $^3\text{H}$  (Table 5). Environmental  $^3\text{H}$  is due primarily to contributions from natural processes and historic nuclear weapons tests. None of the water samples contained a detectable concentration of gross alpha; almost half of the samples contained detectable gross beta concentrations. At these levels, radioactivity in water samples is generally attributed to naturally occurring decay products, primarily from primordial deposits of uranium and thorium, picked up by water as it travels through the earth's crust.

<b>Table 5</b>			
<b>Radionuclide Concentrations in Offsite Water Samples</b>			
<b>(Second Quarter 1996)</b>			
<u>Location</u>	$^3\text{H}$ (pCi/l $\pm$ 2s)	Gross Alpha (pCi/l $\pm$ 2s)	Gross Beta (pCi/l $\pm$ 2s)
<b>Drinking Water</b>			
Aberdeen	20 $\pm$ 90	-4 $\pm$ 5	3 $\pm$ 2
Arco	60 $\pm$ 100	0 $\pm$ 3	0 $\pm$ 2
Atomic City	20 $\pm$ 100	0 $\pm$ 3	3 $\pm$ 2
Blackfoot	160 $\pm$ 100	2 $\pm$ 6	4 $\pm$ 2
Carey	70 $\pm$ 100	2 $\pm$ 4	0 $\pm$ 2
Fort Hall	120 $\pm$ 100	-4 $\pm$ 5	2 $\pm$ 2
Howe	20 $\pm$ 100	-4 $\pm$ 4	0 $\pm$ 2
Minidoka	40 $\pm$ 90	0 $\pm$ 4	3 $\pm$ 2
Monteview	80 $\pm$ 100	0 $\pm$ 40 <sup>a</sup>	8 $\pm$ 3
Mud Lake	10 $\pm$ 100	-2 $\pm$ 2	5 $\pm$ 2
Reno Ranch	40 $\pm$ 100	-1 $\pm$ 4	0 $\pm$ 2
Roberts	-40 $\pm$ 90	-4 $\pm$ 4	2 $\pm$ 2
<i>Roberts Replicate</i>	60 $\pm$ 100	-2 $\pm$ 4	2 $\pm$ 2
Shoshone	40 $\pm$ 100	-2 $\pm$ 4	2 $\pm$ 2
<b>Surface Water</b>			
Alpheus Spring (Twin Falls)	20 $\pm$ 100	0 $\pm$ 5	6 $\pm$ 3
Bill Jones Hatchery (Hagerman)	50 $\pm$ 100	2 $\pm$ 4	2 $\pm$ 2
Clear Spring (Buhl)	40 $\pm$ 100	1 $\pm$ 4	2 $\pm$ 2
<i>Clear Spring Replicate</i>	-20 $\pm$ 100	2 $\pm$ 5	1 $\pm$ 2
Bliss	30 $\pm$ 90	1 $\pm$ 4	2 $\pm$ 2
Idaho Falls	10 $\pm$ 100	-3 $\pm$ 3	1 $\pm$ 2
EPA Maximum Contaminant Level (MCL)	20,000	15	50
<sup>a</sup> Large uncertainty due to high sample turbidity.			

## 4. Milk Sampling

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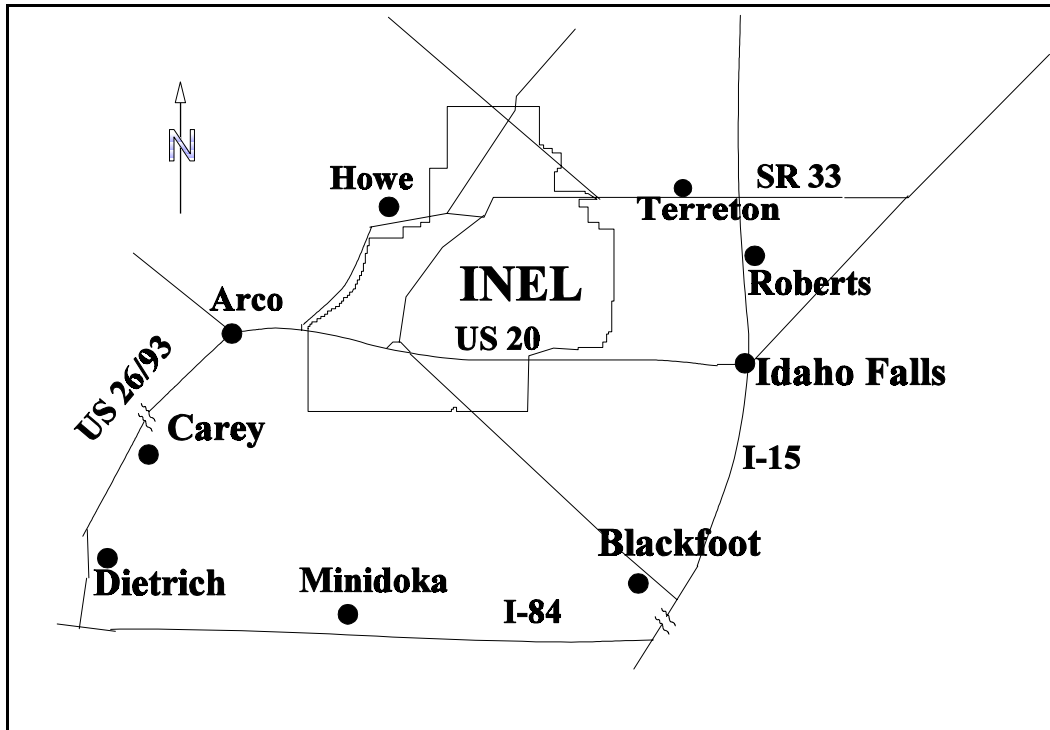


Figure 5 Milk Sampling Locations

## 4. Milk Sampling

### 4.1 Methods

Milk samples were collected weekly in Idaho Falls and monthly at eight other locations around the INEL (Figure 5). Two types of locations were sampled: single family dairies and large commercial dairies. Each milk sample was analyzed for  $^{131}\text{I}$  by placing the sample in a gamma spectrometer calibrated for the  $^{131}\text{I}$  energy peak. Since  $^{131}\text{I}$  has a short half-life (eight days), results are decay-corrected to the time of sample collection. Selected milk samples were also analyzed for  $^3\text{H}$  and  $^{90}\text{Sr}$ .

## 4. Milk Sampling

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### 4.2 Results

A total of 38 milk samples were collected during the second quarter. Iodine-131 was not detected in any of the samples, at a minimum detectable concentration of about  $2 \times 10^{-9}$   $\mu\text{Ci/ml}$ .

Milk samples from four locations (Howe, Roberts, Arco, and Dietrich) were analyzed for  $^3\text{H}$ . One sample (Arco) had a detectable concentration of  $(1.5 \pm 1.0) \times 10^{-7}$   $\mu\text{Ci/ml}$ . Five samples were analyzed for  $^{90}\text{Sr}$  (Table 6). This radionuclide was detected in all samples, four from distant locations and one from a boundary location (Terreton). Most of the distant stations had greater concentrations of  $^{90}\text{Sr}$  than the boundary station. All concentrations were consistent with those reported by the EPA, and are due to worldwide fallout from historic above-ground nuclear weapons tests.

<b>Location</b>	<b>Result</b> <b><math>(\mu\text{Ci/ml} \pm 2s) \times 10^{-9}</math></b>
Blackfoot	$1.8 \pm 0.7$
Carey	$1.7 \pm 0.6$
Idaho Falls	$1.6 \pm 0.6$
Minidoka	$0.9 \pm 0.5$
Terreton	$1.0 \pm 0.7$



## 5. Animal Sampling

### 5.1 Methods

Samples of thyroid, muscle, and liver were taken from sheep grazing on authorized allotments along the eastern side of the INEL in the Tractor Flats area and on the northern portion of the INEL north of TAN (Figure 6). Control samples were taken from the Blackfoot area. In addition, similar organ and tissue samples were taken from road-killed game at locations on or near the INEL. The samples were analyzed for man-made gamma-emitting radionuclides. During this report period, gamma spectrometry protocol was enhanced, as requested by the Foundation, to include detailed analysis of the spectra for a larger number of radionuclides.

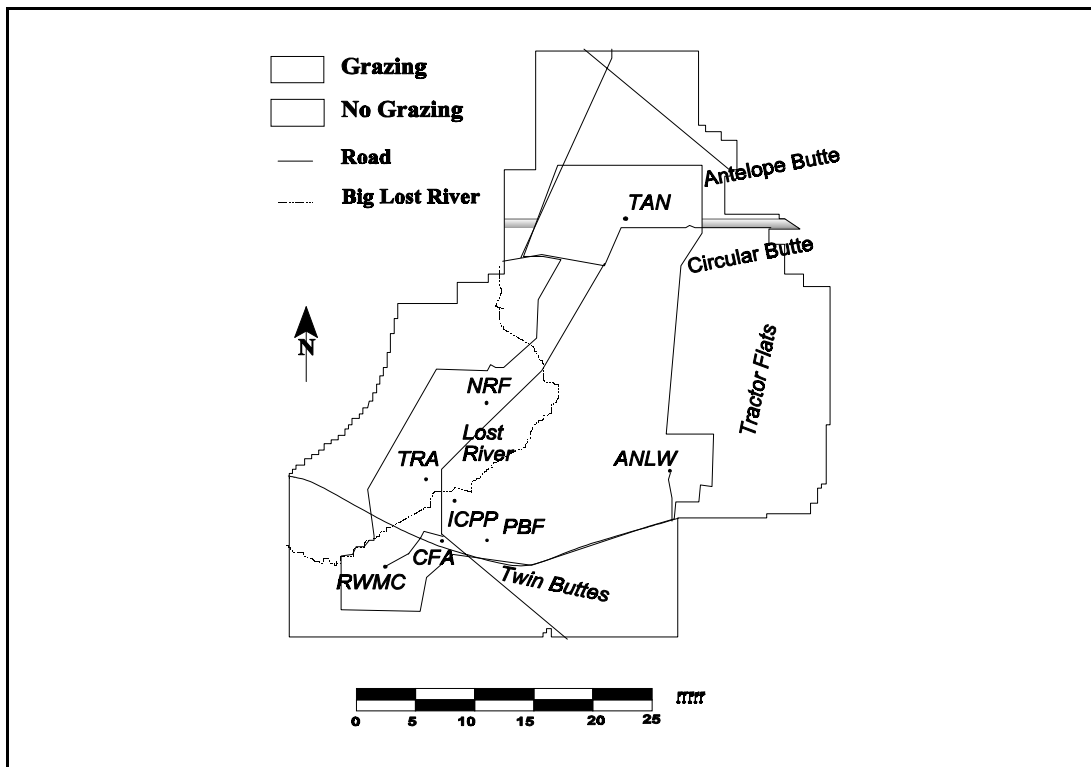


Figure 6 Livestock Grazing Locations on the INEL

## 5. Animal Sampling

### 5.2 Results

Tissue samples from two of the sheep grazing onsite showed detectable concentrations of  $^{137}\text{Cs}$  (Table 7). Neither of the two Blackfoot (control) sheep samples showed detectable concentrations of  $^{137}\text{Cs}$ . The element cesium (Cs) is generally absorbed by all organs and tissues. One of the onsite sheep samples showed detectable concentrations of  $^{60}\text{Co}$ . None of the thyroids from the six sheep indicated detectable concentrations of  $^{131}\text{I}$ .

Five game animals, killed on roads on or near the INEL, were sampled during this report period. The thyroids were analyzed for  $^{131}\text{I}$ . No concentrations were detected. Cesium-137 was detected in four animals; Cobalt-60 was detected in one muscle sample (Table 7).

These radionuclides, whose levels are consistent with past results, are generally attributed to the world-wide inventory of fallout from historic nuclear weapons testing. In some cases, animals which have grazed onsite, particularly game animals whose movements are not controlled, may ingest radioactivity (primarily  $^{137}\text{Cs}$ ) from areas of contaminated soil.

<b>Table 7</b>			
<b>Radionuclide Concentrations in Sheep and Game</b>			
<b>(Second Quarter 1996)</b>			
<u>Location</u>	<u>Tissue/Organ</u>	$^{137}\text{Cs}$ Result ( $\mu\text{Ci/g} \pm 2\text{s}$ ) x $10^{-9}$	$^{60}\text{Co}$ Result ( $\mu\text{Ci/g} \pm 2\text{s}$ ) x $10^{-9}$
<b><u>Sheep</u></b>			
Tractor Flats	Muscle	3.2 ± 2.5	nd <sup>a</sup>
	Liver	nd	2.8 ± 2.1
north of TAN	Muscle	3.0 ± 1.8	nd
	Liver	1.8 ± 1.4	nd
<b><u>Road-killed Game</u></b>			
Highway 26	Muscle (mule deer)	4.9 ± 4.6	nd
Highway 20/26	Muscle (mule deer)	7.3 ± 3.8	3.3 ± 2.9
	Liver (mule deer)	9.1 ± 5.6	nd
Lincoln Blvd.	Muscle (mule deer)	5.6 ± 4.0	nd
Highway 26	Liver (pronghorn)	3.8 ± 3.3	nd

<sup>a</sup> nd = not detected.

## 6. Environmental Radiation

### 6.1 Methods

Environmental radiation is monitored at six boundary and seven distant stations (Figure 7). Environmental radiation is monitored with the use of thermoluminescent dosimeters (TLDs) made of lithium fluoride crystals. The TLDs are placed on posts one meter (3.3 feet) above the ground at field locations and changed every six months in May and November. The crystals detect beta and gamma radiation and store this information in the form of “excited” electrons within the crystals. The TLDs are analyzed by an instrument which heats them under precisely controlled conditions and detects the light they give off. The amount of light is a measure of the amount of environmental radiation.

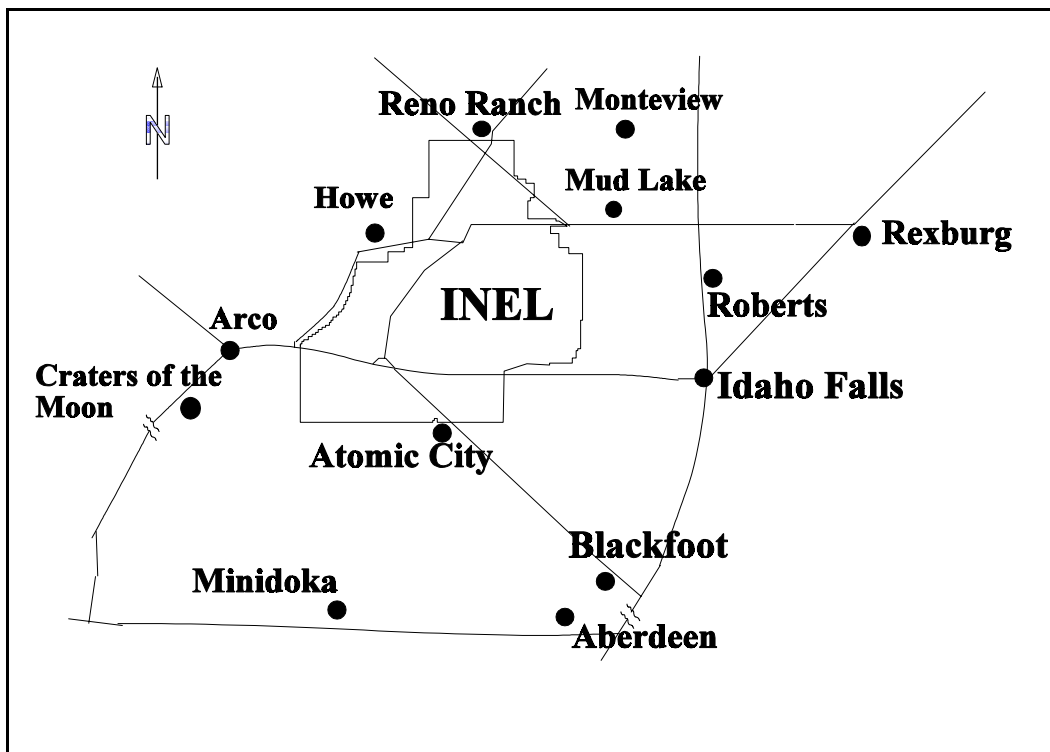


Figure 7 TLD Locations

## 6. Environmental Radiation

### 6.2 Results

The results for the first half of 1996 (November 1995 through April 1996) are shown in Table 8. The results show the exposure levels to be similar to the previous six-month interval, but higher than the same seasonal period a year earlier (November 1994 through April 1995). This is likely due to the relatively light snow cover during the winter of 1995/96. Snow cover shields the dosimeter from radiation exposure from natural radionuclides in the soil. The mean exposure level at the boundary stations was not statistically greater than the mean exposure level at the distant stations.

<b>Table 8</b>			
<b>Environmental Radiation Exposure (mR) for Nov. 1995-May 1996</b>			
<b>Location</b>	<b>11/94-5/95 Exposure (mR ± 2s)</b>	<b>5/95-11/95 Exposure (mR ± 2s)</b>	<b>11/95-5/96 Exposure (mR ± 2s)</b>
<b>Distant Locations</b>			
Aberdeen	49 ± 2	59 ± 2	56 ± 4
Blackfoot	54 ± 3	63 ± 2	57 ± 3
Craters of the Moon	52 ± 2	62 ± 3	55 ± 2
Idaho Falls	57 ± 4	63 ± 3	61 ± 3
Minidoka	51 ± 1	54 ± 2	56 ± 4
Rexburg	50 ± 1	59 ± 3	60 ± 3
Roberts	59 ± 3	59 ± 4	71 ± 5
<b>Group Mean<sup>a</sup></b>	<b>53 ± 3</b>	<b>60 ± 3</b>	<b>59 ± 5</b>
<b>Boundary Locations</b>			
Arco	55 ± 2	63 ± 2	58 ± 3
Atomic City	59 ± 2	65 ± 4	70 ± 5
Howe	54 ± 3	58 ± 3	57 ± 5
Monteview	56 ± 3	62 ± 3	60 ± 2
Mud Lake	59 ± 3	58 ± 6	62 ± 2
Reno Ranch	55 ± 2	58 ± 3	57 ± 3
<b>Group Mean<sup>a</sup></b>	<b>56 ± 2</b>	<b>61 ± 3</b>	<b>61 ± 5</b>
a. Mean ± 95% confidence interval.			

## **Appendix**

### **Weekly Gross Alpha and Gross Beta Concentrations in Air**

**Table A-1**  
**Weekly Gross Alpha Concentrations in Air**

<u>Location</u>	<u>Weekly Collection Date</u>	<u>Concentration ± 2s Uncertainty (10<sup>-15</sup> μCi/ml)</u>
	<u>Distant Locations</u>	
Blackfoot	04/10	2.0 ± 0.8
	04/17	1.2 ± 0.7
	04/24	1.5 ± 0.7
	05/01	0.7 ± 0.6
	05/08	3.1 ± 1.0
	05/15	1.9 ± 0.8
	05/22	1.2 ± 0.7
	05/29	1.0 ± 0.7
	06/05	3.0 ± 0.9
	06/12	1.9 ± 0.9
	06/19	1.6 ± 0.8
	06/26	1.7 ± 0.8
	07/03	2.2 ± 0.9
Craters of the Moon	04/10	0.7 ± 0.7
	04/17	-0.2 ± 0.5
	04/24	0.4 ± 0.6
	05/01	0.9 ± 0.8
	05/08	2.5 ± 1.0
	05/15	1.2 ± 0.8
	05/22	1.0 ± 0.8
	05/29	1.1 ± 0.8
	06/05	1.8 ± 0.9
	06/12	1.2 ± 0.8
	06/19	1.8 ± 0.9
	06/26	1.1 ± 0.8
	07/03	2.0 ± 0.9
Idaho Falls	04/10	2.2 ± 0.8
	04/17	1.3 ± 0.7
	04/24	1.2 ± 0.7
	05/01	1.8 ± 0.8
	05/08	1.4 ± 0.8
	05/15	1.1 ± 0.7
	05/22	1.5 ± 0.8
	05/29	0.5 ± 0.6
	06/05	1.1 ± 0.8
	06/12	2.1 ± 0.9
06/19	1.4 ± 0.8	

**Table A-1 (Cont.)  
Weekly Gross Alpha Concentrations in Air**

<u>Location</u>	<u>Weekly Collection Date</u>	<u>Concentration ± 2s Uncertainty (10<sup>-15</sup> μCi/ml)</u>
	06/26	1.4 ± 0.8
	07/03	1.6 ± 0.9
Rexburg	04/10	1.1 ± 0.7
	04/17	0.9 ± 0.6
	04/24	0.9 ± 0.6
	05/01	1.8 ± 0.8
	05/08	2.4 ± 0.9
	05/15	1.3 ± 0.7
	05/22	0.7 ± 0.6
	05/29	0.9 ± 0.6
	06/05	2.4 ± 0.9
	06/12	2.0 ± 0.8
	06/19	1.2 ± 0.7
	06/26	2.0 ± 0.8
	07/03	2.0 ± 0.9
<b>Boundary Locations</b>		
Arco	04/10	0.8 ± 0.6
	04/17	0.8 ± 0.6
	04/24	0.2 ± 0.5
	05/01	0.2 ± 0.5
	05/08	2.2 ± 0.9
	05/15	0.9 ± 0.7
	05/22	1.1 ± 0.7
	05/29	0.6 ± 0.6
	06/05	1.5 ± 0.7
	06/12	1.3 ± 0.7
	06/19	0.8 ± 0.7
	06/26	1.0 ± 0.7
	07/03	1.1 ± 0.8
Atomic City	04/10	1.1 ± 0.7
	04/17	0.7 ± 0.6
	04/24	0.6 ± 0.6
	05/01	0.5 ± 0.6
	05/08	1.4 ± 0.8
	05/15	1.2 ± 0.7
	05/22	0.7 ± 0.6
	05/29	0.7 ± 0.6
	06/05	1.4 ± 0.7

**Table A-1 (Cont.)  
Weekly Gross Alpha Concentrations in Air**

<u>Location</u>	<u>Weekly Collection Date</u>	<u>Concentration ± 2s Uncertainty (10<sup>-15</sup> μCi/ml)</u>
	06/12	1.8 ± 0.8
	06/19	1.3 ± 0.7
	06/26	0.7 ± 0.7
	07/03	1.3 ± 0.7
FAA Tower	04/10	0.8 ± 0.7
	04/17	0.2 ± 0.6
	04/24	0.7 ± 0.7
	05/01	1.1 ± 0.8
	05/08	1.9 ± 0.9
	05/15	1.7 ± 0.8
	05/22	0.7 ± 0.7
	05/29	0.9 ± 0.7
	06/05	1.1 ± 0.8
	06/12	1.4 ± 0.8
	06/19	2.0 ± 0.9
	06/26	1.3 ± 0.8
	07/03	1.6 ± 0.9
Howe	04/10	1.2 ± 0.7
	04/17	0.3 ± 0.6
	04/24	0.5 ± 0.6
	05/01	1.1 ± 0.8
	05/08	2.6 ± 1.0
	05/15	1.7 ± 0.8
	05/22	0.4 ± 0.6
	05/29	0.3 ± 0.5
	06/05	1.1 ± 0.8
	06/12	1.2 ± 0.8
	06/19	0.8 ± 0.7
	06/26	-0.5 ± 0.3
	07/03	2.2 ± 0.9
Monteview	04/10	1.7 ± 0.8
	04/17	0.4 ± 0.6
	04/24	1.1 ± 0.7
	05/01	1.5 ± 0.8
	05/08	2.8 ± 1.0



**Table A-1 (Cont.)  
Weekly Gross Alpha Concentrations in Air**

<u>Location</u>	<u>Weekly Collection Date</u>	<u>Concentration ± 2s Uncertainty (10<sup>-15</sup> μCi/ml)</u>
	05/15	1.2 ± 0.8
	05/22	0.1 ± 0.5
	05/29	0.3 ± 0.6
	06/05	0.8 ± 0.7
	06/12	2.3 ± 0.9
	06/19	1.1 ± 0.7
	06/26	1.2 ± 0.8
	07/03	1.7 ± 0.8
Mud Lake	04/10	1.7 ± 0.8 (1.9 ± 0.8)
(Replicate)	04/17	0.6 ± 0.7 (0.2 ± 0.6)
	04/24	0.6 ± 0.6 (0.5 ± 0.6)
	05/01	2.1 ± 1.0 (2.1 ± 1.0)
	05/08	2.1 ± 0.9 (3.4 ± 1.1)
	05/15	2.4 ± 0.9 (1.5 ± 0.9)
	05/22	0.9 ± 0.7 (0.7 ± 0.7)
	05/29	0.4 ± 0.6 (0.7 ± 0.7)
	06/05	2.1 ± 0.8 (1.7 ± 0.8)
	06/12	1.5 ± 0.8 (1.9 ± 0.9)
	06/19	1.6 ± 0.8 (1.8 ± 0.9)
	06/26	1.2 ± 0.8 (1.8 ± 0.9)
	07/03	1.8 ± 0.8 (1.5 ± 0.8)
Reno Ranch	04/10	0.7 ± 0.7
	04/17	0.5 ± 0.6
	04/24	0.2 ± 0.5
	05/01	1.1 ± 0.8
	05/08	1.4 ± 0.8
	05/15	1.1 ± 0.7
	05/22	0.4 ± 0.6
	05/29	0.2 ± 0.6
	06/05	1.0 ± 0.7
	06/12	1.7 ± 0.9
	06/19	1.1 ± 0.7
	06/26	0.9 ± 0.7
	07/03	1.2 ± 0.8
INEL Locations		
EFS	04/10	0.7 ± 0.6
	04/17	0.7 ± 0.6
	04/24	0.1 ± 0.5

**Table A-1 (Cont.)**  
**Weekly Gross Alpha Concentrations in Air**

<u>Location</u>	<u>Weekly Collection Date</u>	<u>Concentration ± 2s Uncertainty (10<sup>-15</sup> μCi/ml)</u>
	05/01	1.2 ± 0.8
	05/08	1.8 ± 0.9
	05/15	1.2 ± 0.7
	05/22	0.4 ± 0.6
	05/29	0.5 ± 0.6
	06/05	1.6 ± 0.9
	06/12	1.2 ± 0.8
	06/19	1.5 ± 0.8
	06/26	1.3 ± 0.8
	07/03	1.7 ± 0.9
Main Gate (Replicate)	04/10	1.6 ± 0.8 (0.8 ± 0.7)
	04/17	0.6 ± 0.6 (0.4 ± 0.7)
	04/24	0.1 ± 0.6 (0.1 ± 0.6)
	05/01	1.2 ± 0.8 (0.5 ± 0.7)
	05/08	2.3 ± 1.0 (2.0 ± 1.0)
	05/15	1.1 ± 0.8 (1.4 ± 0.8)
	05/22	0.0 ± 0.6 (0.2 ± 0.7)
	05/29	-0.6 ± 0.4 (0.7 ± 0.7)
	06/05	1.6 ± 0.9 (1.5 ± 0.8)
	06/12	2.1 ± 0.9 (1.4 ± 0.9)
	06/19	1.9 ± 0.9 (1.2 ± 0.8)
	06/26	1.5 ± 0.9 (0.6 ± 0.7)
	07/03	2.7 ± 1.0 (0.8 ± 1.3)
Van Buren	04/10	1.3 ± 0.8
	04/17	0.2 ± 0.6
	04/24	0.6 ± 0.6
	05/01	0.6 ± 0.7
	05/08	1.8 ± 0.9
	05/15	1.3 ± 0.8
	05/22	0.9 ± 0.7
	05/29	0.2 ± 0.6
	06/05	1.9 ± 0.9
	06/12	1.7 ± 0.9
	06/19	1.3 ± 0.8
	06/26	1.3 ± 0.8
	07/03	0.8 ± 0.7

**Table A-2**  
**Weekly Gross Beta Concentrations in Air**

<u>Location</u>	<u>Weekly Collection Date</u>	<u>Concentration ± 2s Uncertainty (10<sup>-15</sup> μCi/ml)</u>
	Distant Locations	
Blackfoot	04/10	22 ± 4
	04/17	13 ± 4
	04/24	13 ± 4
	05/01	8 ± 3
	05/08	26 ± 4
	05/15	20 ± 4
	05/22	11 ± 4
	05/29	8 ± 4
	06/05	26 ± 4
	06/12	27 ± 5
	06/19	25 ± 4
	06/26	20 ± 5
	07/03	20 ± 4
Craters of the Moon	04/10	22 ± 5
	04/17	18 ± 4
	04/24	11 ± 4
	05/01	22 ± 5
	05/08	30 ± 5
	05/15	20 ± 5
	05/22	16 ± 5
	05/29	17 ± 5
	06/05	24 ± 5
	06/12	27 ± 5
	06/19	38 ± 5
	06/26	26 ± 5
	07/03	26 ± 5
Idaho Falls	04/10	16 ± 4
	04/17	12 ± 4
	04/24	12 ± 4
	05/01	15 ± 4
	05/08	21 ± 5
	05/15	17 ± 4
	05/22	13 ± 4
	05/29	10 ± 4
	06/05	13 ± 4
	06/12	18 ± 5

**Table A-2 (Cont.)  
Weekly Gross Beta Concentrations in Air**

<u>Location</u>	<u>Weekly Collection Date</u>	<u>Concentration ± 2s Uncertainty (10<sup>-15</sup> μCi/ml)</u>
	06/19	22 ± 5
	06/26	14 ± 5
	07/03	19 ± 5
Rexburg	04/10	22 ± 4
	04/17	17 ± 4
	04/24	11 ± 4
	05/01	19 ± 4
	05/08	25 ± 4
	05/15	22 ± 4
	05/22	12 ± 4
	05/29	9 ± 4
	06/05	23 ± 4
	06/12	28 ± 4
	06/19	32 ± 5
	06/26	15 ± 4
	07/03	21 ± 5
<b>Boundary Locations</b>		
Arco	04/10	19 ± 4
	04/17	11 ± 4
	04/24	10 ± 3
	05/01	17 ± 4
	05/08	26 ± 5
	05/15	19 ± 4
	05/22	10 ± 4
	05/29	14 ± 4
	06/05	24 ± 4
	06/12	23 ± 4
	06/19	25 ± 5
	06/26	15 ± 4
	07/03	21 ± 5
Atomic City	04/10	26 ± 4
	04/17	18 ± 4
	04/24	15 ± 4
	05/01	16 ± 4
	05/08	29 ± 5
	05/15	18 ± 4
	05/22	10 ± 4
	05/29	6 ± 3

**Table A-2 (Cont.)  
Weekly Gross Beta Concentrations in Air**

<u>Location</u>	<u>Weekly Collection Date</u>	<u>Concentration ± 2s Uncertainty (10<sup>-15</sup> μCi/ml)</u>
	06/05	23 ± 4
	06/12	29 ± 5
	06/19	30 ± 5
	06/26	17 ± 4
	07/03	20 ± 4
FAA	04/10	23 ± 5
Tower	04/17	10 ± 4
	04/24	11 ± 4
	05/01	13 ± 4
	05/08	26 ± 5
	05/15	15 ± 4
	05/22	13 ± 4
	05/29	8 ± 4
	06/05	21 ± 5
	06/12	21 ± 5
	06/19	30 ± 5
	06/26	23 ± 5
	07/03	24 ± 5
Howe	04/10	25 ± 4
	04/17	13 ± 4
	04/24	11 ± 4
	05/01	19 ± 5
	05/08	27 ± 5
	05/15	18 ± 4
	05/22	17 ± 4
	05/29	16 ± 4
	06/05	23 ± 5
	06/12	25 ± 5
	06/19	24 ± 5
	06/26	21 ± 4
	07/03	20 ± 4
Monteview	04/10	21 ± 4
	04/17	15 ± 4
	04/24	11 ± 4
	05/01	22 ± 5
	05/08	22 ± 5
	05/15	15 ± 5

**Table A-2 (Cont.)  
Weekly Gross Beta Concentrations in Air**

<u>Location</u>	<u>Weekly Collection Date</u>	<u>Concentration ± 2s Uncertainty (10<sup>-15</sup> μCi/ml)</u>
	05/22	10 ± 4
	05/29	9 ± 4
	06/05	23 ± 5
	06/12	25 ± 5
	06/19	26 ± 5
	06/26	12 ± 5
	07/03	21 ± 5
Mud Lake	04/10	31 ± 5 (27 ± 5)
(Replicate)	04/17	14 ± 4 (13 ± 4)
	04/24	14 ± 4 (13 ± 4)
	05/01	16 ± 5 (23 ± 6)
	05/08	28 ± 5 (31 ± 6)
	05/15	18 ± 4 (21 ± 5)
	05/22	13 ± 4 (12 ± 4)
	05/29	10 ± 4 (13 ± 4)
	06/05	27 ± 5 (27 ± 5)
	06/12	26 ± 5 (26 ± 5)
	06/19	32 ± 5 (35 ± 5)
	06/26	18 ± 4 (25 ± 3)
	07/03	24 ± 5 (31 ± 5)
Reno	04/10	22 ± 4
Ranch	04/17	13 ± 4
	04/24	9 ± 4
	05/01	15 ± 4
	05/08	25 ± 5
	05/15	19 ± 4
	05/22	12 ± 4
	05/29	14 ± 4
	06/05	24 ± 5
	06/12	26 ± 5
	06/19	27 ± 5
	06/26	19 ± 5
	07/03	22 ± 5
INEL Locations		
EFS	04/10	26 ± 4
	04/17	16 ± 4
	04/24	13 ± 4
	05/01	19 ± 5

**Table A-2 (Cont.)  
Weekly Gross Beta Concentrations in Air**

<u>Location</u>	<u>Weekly Collection Date</u>	<u>Concentration ± 2s Uncertainty (10<sup>-15</sup> μCi/ml)</u>
	05/08	31 ± 5
	05/15	21 ± 4
	05/22	14 ± 4
	05/29	11 ± 4
	06/05	27 ± 5
	06/12	27 ± 5
	06/19	24 ± 5
	06/26	17 ± 4
	07/03	23 ± 5
Main Gate	04/10	26 ± 4 (22 ± 5)
(Replicate)	04/17	11 ± 4 (12 ± 5)
	04/24	12 ± 4 (12 ± 5)
	05/01	16 ± 4 (16 ± 5)
	05/08	29 ± 6 (28 ± 5)
	05/15	28 ± 5 (21 ± 5)
	05/22	15 ± 4 (16 ± 5)
	05/29	17 ± 4 (14 ± 4)
	06/05	26 ± 5 (19 ± 5)
	06/12	33 ± 5 (25 ± 5)
	06/19	29 ± 5 (27 ± 5)
	06/26	26 ± 5 (19 ± 5)
	07/03	30 ± 5 (21 ± 9)
Van Buren	04/10	24 ± 4
	04/17	11 ± 4
	04/24	10 ± 4
	05/01	16 ± 4
	05/08	22 ± 5
	05/15	18 ± 4
	05/22	7 ± 4
	05/29	9 ± 4
	06/05	21 ± 5
	06/12	23 ± 5
	06/19	28 ± 5
	06/26	21 ± 4
	07/03	19 ± 5

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